

DESCRIPTION

HUMAN IgM ANTIBODY LYSING ACTIVATED LYMPHOCYTES UNDER MEDIATION BY HOMOLOGOUS COMPLEMENT

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Technical Field

The invention relates to a human IgM monoclonal antibody reactive to HIV-infected cells that lyses activated lymphocytes by reacting with a differentiated antigen of the activated lymphocytes under medication by homologous human complements, and a remedy for autoimmune diseases containing the monoclonal antibody.

Background Art

15 Various immunosuppressants such as cyclosporin and FK506 have been developed for controlling biological immunoreactions in collagen diseases, autoimmune diseases and rejection for organ transplantation. However, since such immunosuppressants are reactive to cells other than immunocompetent cells, side effects of these agent should be taken into consideration.

20 Various methods have been investigated for using antibodies that specifically react with target cells. For example, the target cell with which the antibody has reacted is expected to be lysed by reacting with a complement. While there are species-specific complement control membrane

factors (such as DAF, decay accelerating factor; MCP, membrane cofactor protein; and HRF20, 20 kDa homologous restriction factor) on human cell membranes, they can induce no cytolysis reaction via complement reactions for

5 preventing reactions among homologous human complements.

On the other hand, it was found that IgM antibodies in human serum that react with the HIV-infected cells are able to yield the cytolysis reaction of the HIV-infected cells via the human complement by overcoming the complement

10 control membrane factors. It was revealed that the IgM antibody can exhibit such action as described above against gangliosides such as GM2 and Gg4 whose expression is enhanced by HIV-infection (Japanese Patent Application Laid-Open No. 9-227409).

15 L55 has been reported as the human IgM monoclonal antibody against GM2 of the gangliosides, wherein L55 is produced by immortalizing human B lymphoblast strain with EB virus. The HIV-infected cells after treating with this human IgM monoclonal antibody have been found to yield
20 cytolysis via a reaction with the human complement.

Disclosure of Invention

An object of the invention is to provides a remedy for controlling immunological response containing a human IgM
25 monoclonal antibody that specifically reacts with activated lymphocytes to induce cytolysis under mediation by

homologous complements.

As a result of intensive studies for solving the problems above, the invention provides a human IgM monoclonal antibody reactive to HIV-infected cells that lyse
5 activated human lymphocytes under mediation by homologous human complements.

The invention for solving the problems above provides an HIV remedy that cures transplantation rejection response and autoimmune diseases caused by excessive response of T-
10 lymphocytes as well as HIV infection diseases by eliminating activated lymphocytes by cytolyses using a human IgM monoclonal antibody that reacts with HIV-infected cells and activated lymphocytes.

More specifically, the invention for solving the
15 problems above provides the human IgM monoclonal antibody according to any one of first to second aspects, wherein the human IgM monoclonal antibody that reacts with the activated lymphocytes and the HIV-infected cells is 9F11 antibody having a base sequence of the H-chain variable region
20 represented by sequence number 1.

More specifically, the invention for solving the problems above provides the human IgM monoclonal antibody according to any one of first to third aspects, wherein the human IgM monoclonal antibody that reacts with the activated
25 lymphocytes and the HIV-infected cells is 9F11 antibody having a base sequence of the L-chain variable region

represented by sequence number 2.

Brief Description of the Drawings

5 Fig. 1 shows specificity of 9F11 antibody.

The result of flow cytometry analysis shows that HIV-infected cells are stained with 9F11 antibodies while non-infected cells are not.

10 Fig. 2 shows specificity of 9F11 antibody to peripheral blood lymphocytes.

The drawing shows that 9F11 antibody reacts with the activated lymphocytes stimulated with PHA while it does not react with normal peripheral blood lymphocytes (PBMC: peripheral blood lymphocyte).

15 Fig. 3 shows a cell impairing reaction by 9F11 antibody under mediation by the complement.

(A) Almost all cells are destroyed 4 hours after addition of 2 µg/ml of 9F11 antibody and fresh human serum (including a complement component) to HIV-1 infected cells
20 MOLT-4/IIIB. No effects have been observed at all when the serum is not added or on non-infected cells MOLT-4.

(B) 9F11 antigen is induced by activating the peripheral blood lymphocytes using PHA, and the cells are impaired by 9F11 antibody and the complement as in the HIV-1 infected
25 cells (FHS: fresh human serum (used as a complement source), PHA: human lymphocyte activating agent, %⁵¹Cr release =

mortality of cells, PBMC: peripheral blood lymphocyte).

Fig. 4 schematically illustrates 9F11 μ -chain expression plasmid construct.

5 Best Mode for Carrying Out the Invention

While the invention is described in detail with reference to examples, the technical scope of the invention is by no means restricted to these examples.

For solving the problems above, the inventors of the
10 invention immunized HIV-infected cells of a mouse (TC mouse: trans-chromosome mouse; prepared by Kirin Brewery Co., Ltd.) into which human immunoglobulin gene-containing chromosomes had been introduced, and obtained a mouse that produces human antibodies that specifically react with HIV-infected
15 cells. Hybridoma was prepared by a conventional method by fusing spleen cells of the immunized mouse with a mouse myeloma cell strain. Clones producing the monoclonal antibody that react with the HIV-infected cells and lead the infected cells to cytolysis in the presence of a human
20 complement were selected from the hybridoma. The selected hybridomas were named as 9F11 cell strain. A 9F11 antibody that is a monoclonal antibody produced by 9F11 cell strain is a human IgM monoclonal antibody comprising human μ -chain and human κ -chain. While 9F11 antibody was able to yield
25 cytolysis by reacting with the HIV-infected cells under mediation by the human complement, the antibody also caused

cytolysis for activated lymphocytes of non-infected lymphocytes. Accordingly, the response to the HIV-infected cells may be comprehended that the lymphocyte is brought to a certain kind of activated state by HIV infection, and the
5 HIV-infected cells are lysed under mediation by the human complement by expressing an antigen (9F11 antigen) against 9F11 as a differentiated antigen. In other words, 9F11 antigen is a differentiated antigen that is expressed by activation of the lymphocyte, and 9F11 antibody that induces
10 cytolysis under mediation by the complement by reacting with the antigen specifically leads the activated lymphocyte to cytolysis under mediation by the complement. It was therefore made clear that a remedy containing 9F11 antibody can be utilized in the treatment for suppressing the
15 activated lymphocyte. The invention has been completed based on the discoveries described above. The cell strain 9F11 that produces 9F11 antibodies of the invention was deposited with National Institute of Advanced Industrial Science and Technology, International Patent Organism
20 Depository (Chuo-6, Higashi 1-1-1, Tsukuba City, Ibaraki Pref.), on May 8, 2003, with an accession number of FERM BP-8379.

Table 1 shows the results of base sequence analysis of the genes in the variable regions in κ -chain and μ -chain,
25 respectively, encoding 9F11 antibody. The base sequence of the constant region is approximately the same as the base

sequence of reported genes.

TABLE 1

Base Sequence of μ -Chain Variable Region

5 **GCTGAATTCTGGCTGACCAGGGCAGTCACCAGAGCTCCAGACAATGTCTG**
 TCTCCTTCCTCATCTTCCTGCCCCGTGCTGGGCCTCCCATGGGGTGTCTGT
 CACAGGTACAGCTGCAGCAGTCAGGTCCAGGACTGGTGAAGCCCCGCGCAG
 ACCCTCTCACTCACCTGTGCCATCTCCGGGGACAGTGTCTCTAGCAACAGT
 GCTACTTGGAAGTGGATCAGGCAGTCCCCATTGAGAGGCCTTGAGTGGCT
10 **GGGAAGGACATACTACAGGTCCAAGTGGTATAATGATTATGCAGTATCTG**
 TGAAAAGTCGAATAACCATCAACCCAGACACATCCAAGAACCAGTTCTCCC
 TGCAGCTGAACTCTGTGACTCCCGAGGACACGGCTGTGTATTACTGTGCA
 AGAGAGAATTACTATGGTTCGGGGAGGTACAAGTGGTTCGACCCCTGGGG
 CCAGGGAACCCTGGTCACCGTCTCCTCA

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Base Sequence of κ -Chain Variable Region

TGTCAGGACACAGCATGGACATGAGGGTCCCCGCTCAGCTCCTGGGGCTC
 CTGCTGCTCTGGTTCCCAGGTTCCAGATGCGACATCCAGATGACCCAGTC
 TCCATCTTCCGTGTCTGCATCTGTAGGAGACAGAGTCACCATCACTTGTCG
20 **GGCGAGTCAGGGTATTAGCAGCTGGTTAGCCTGGTATCAGCAGAAACCAG**
 GGAAAGCCCCTAAGCTCCTGATCTATGATGCATCCAGTTTGCAAAGTGGG
 GTCCCATCAAGGTTCAAGCGGCAGTGGATCTGGGACAGATTTCCTCTCAC
 CATCAGCAGCCTGCAGCCTGAAGATTTTGCAACTTACTATTGTCAACAGGC
 TAACAGTTTCCCTCTCACTTTCGGCGGAGGGACCAAGGTGGAGATCAAA

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The composition of the invention, which is used for an immunological reaction control agent containing the human IgM monoclonal antibody that specifically reacts with the

activated lymphocytes and induces cytolysis under mediation by the homologous complement, may be obtained by combining with physiologically acceptable carriers. The physiologically acceptable carrier is known in the art, and includes physiological buffered saline or other aqueous solutions having a buffer action, or solvents such as glycols, glycerol, oils (for example olive oil) or injectable organic esters. The physiologically acceptable carrier may include compounds that stabilize human IgM antibody or enhance absorption thereof. Examples of the physiologically acceptable compounds include sugars such as glucose, sucrose and dextran; antioxidants such as ascorbic acid and glutathione; chelating agents; proteins such as albumin; and other stabilizers and excipients. Various immunosuppressants such as cyclosporin and FK506 as well as other immunosuppressants may be added to the composition. Any combinations of the physiologically acceptable carriers may be selected depending on administration path and disease of object.

20

Example 1. Specificity of 9F11 antibody

Test cells were suspended in a culture medium in a concentration of 1×10^6 cells/ml, and an equal volume of a 9F11 antibody solution in a concentration of 10 $\mu\text{g/ml}$ was added to the suspension followed by reacting for 30 minutes. The test cells were washed, linked 9F11 was stained with

fluorescence-labeled antihuman IgM antibody, and the cells were analyzed by flow cytometry. The results showed that 9F11 antigen is expressed by HIV infection since, although MOLT-4 cells and CEM cells as human cell strains were not stained, the cells infected with HIV-1 IIIB strain and MN strain were strongly stained (Fig. 1). Since HIV infection induces activation of the lymphocytes, staining of the activated lymphocytes was investigated after cultivating for 3 days by adding phytohemagglutinin (PHA) to the lymphocyte in the peripheral blood cells of a normal person. It was made clear that 9F11 antigen is a differentiated antigen expressed in the activated T-lymphocytes (Fig. 2), since the activated T-lymphocytes stimulated with PHA was strongly stained, although no staining was observed in the peripheral blood cells and peripheral blood lymphocytes not stimulated.

Example 2. Cell impairing reaction by 9F11 antibody under mediation by complement

Test cells were labeled with ^{51}Cr as a radioactive isotope in advance, and 40 μl of 9F11 solutions in various concentrations and 20 μl of human fresh serum (complement serum) were added to the labeled test cells (suspended in a culture medium at a concentration of $5 \times 10^5/\text{ml}$) to allow the mixture to react for 4 hours on a microtiter plate. After the reaction, the cells were precipitated by centrifugation of the plate, and the radioactivity of ^{51}Cr released in the

supernatant by cytolysis was measured as an index of the cytolysis reaction. The cells expressing 9F11 antigen such as MOLT-4/IIIB (MOLT-4 cells infected with HIV-1) and peripheral blood derived lymphoblasts activated with PHA
5 were lysed under mediation by the complement in the presence of 2 µg/ml of 9F11. On the contrary, no cytolysis was observed by using an inactivated serum after heating human serum at 56°C or C9 deficient human fresh serum. Since cytolysis occurs by adding purified C9 to the C9 deficient
10 human fresh serum, it was concluded that the human complement reaction is essential for the cytolysis reaction (Fig. 3).

Example 3. Reconstruction of antibody by gene engineering

An example of the method for reconstruction of 9F11
15 antibody based on the base sequence of the variable region of 9F11 antibody shown in TABLE 1 will be described below, wherein 9F11 antibody-producing cell strains were established using gene engineering such as a shot-gun ligation method (Grundstrom, T. et al., Nucleic Acid Res. 13,
20 3305-3316, 1985).

Amino acid sequences of the variable region of 9F11 antibody were obtained by translating the base sequence in the table. There are many base sequences encoding the amino acid sequences in the variable region of 9F11 antibody as
25 shown in Table 2, such as the base sequence of the variable region of original 9F11 antibody as well as those obtained

by changing used codons. Base sequences having certain kinds of restriction enzyme recognition fragment were selected from the sequences for every length capable of chemically synthesizing as oligonucleotides (Table 2).

TABLE 2: Examples of cDNA encoding equivalent amino acids in the amino acid sequences of 9F11 antibody

1	M	S	V	S	F	L	I	F	L	P	V	L	G	L	P	W	G	V	L	S
	ATA	TCT	GTT	TCT	TTT	TTA	ATT	TTT	TTA	CCT	GTT	TTA	GGT	TTA	CCT	TGA	GGT	GTT	TTA	TCT
	ATG	TCC	GTC	TCC	TTC	TTG	ATC	TTC	TTG	CCC	GTC	TTG	GGC	TTG	CCC	TGG	GGC	GTC	TTG	TCC
	TCA	GTA	TCA		CTT				CTT	CCA	GTA	CTT	GGT	CTT	CCA		GGT	GTA	CTT	TCA
	TCG	GTG	TCG		CTC				CTC	CCG	GTG	CTC	GGG	CTC	CCG		GGG	GTG	CTC	TCG
	AGT		AGT		CTA				CTA			CTA		CTA					CTA	AGT
	AGC		AGC		CTG				CTG			CTG		CTG					CTG	AGC

21

Q V Q L Q Q S G P G L V K P A Q T L S L
 CAA GTT CAA TTA CAA CAA TCT GGT CCT GGT TTA GTT AAA CCT GCT CAA ACT TTA TCT TTA
 CAG GTC CAG TTG CAG CAG TCC GGC CCC GGC TTG GTC AAG CCC GCC CAG ACC TTG TCC TTG
 GTA CTT TCA GGA CCA GGA CTT GTA CCA GCA ACA CTT TCA CTT
 GTG CTC TCG GGG CCG GGG CTC GTG CCG GCG ACG CTC TCG CTC
 CTA CTA AGT CTA CTA AGT CTA
 CTG CTG AGC CTG CTG AGC CTG

- 13 -

41

T C A I S G D S V S S N S A T W N W I R
 ACT TGT GCT ATT TCT GGT GAT TCT GTT TCT TCT AAT TCT GCT ACT TGA AAT TGA ATT CGT
 ACC TGC GCC ATC TCC GGC GAC TCC GTC TCC TCC AAC TCC GCC ACC TGG AAC TGG ATC CGC
 ACA GCA TCA GGA TCA GTA TCA TCA TCA GCA ACA CGA
 ACG GCG TCG GGG TCG GTG TCG TCG TCG GCG ACG CGG
 AGT AGT AGT AGT AGT AGT
 AGC AGC AGC AGC AGC AGC

61

Q	S	P	L	R	G	L	E	W	L	G	R	T	Y	Y	R	S	K	W	Y										
CAA TCT CCT TTA CGT GGT TTA GAA TGA TTA GGT CGT ACT TAT TAT CGT TCT AAA TGA TAT																													
CAG TCC CCC TTG CGC GGC TTG GAG TGG TTG GGC CGC ACC TAC TAC CGC TCC AAG TGG TAC																													
TCA CCA CTT CGA GGA CTT										CTT GGA CGA ACA										CGA TCA									
TCG CCG CTC CGG GGG CTC										CTC GGG CGG ACG										CGG TCG									
AGT					CTA					CTA					CTA					AGT									
AGC					CTG					CTG					CTG					AGC									

81

N D Y A V S V K S R I T I N P D T S K N
AAT GAT TAT GCT GTT TCT GTT AAA TCT CGT ATT ACT ATT AAT CCT GAT ACT TCT AAA AAT
AAC GAC TAC GCC GTC TCC GTC AAG TCC CGC ATC ACC AAC CCC GAC ACC TCC AAG AAC
GCA GTA TCA GTA TCA CGA ACA CCA ACA TCA
GCG GTG TCG GTG TCG CGG ACG CCG ACG TCG
AGT AGT AGT
AGC AGC AGC

101

Q F S L Q L N S V T P E D T A V Y Y C A
CAA TTT TCT TTA CAA TTA AAT TCT GTT ACT CCT GAA GAT ACT GCT GTT TAT TAT TGT GCT
CAG TTC TCC TTG CAG TTG AAC TCC GTC ACC CCC GAG GAC ACC GCC GTC TAC TAC TGC GCC
TCA CTT CTT TCA GTA ACA CCA ACA GCA GTA GCA
TCG CTC CTC TCG GTG ACG CCG ACG GCG GTG GCG
AGT CTA CTA AGT
AGC CTG CTG AGC

R E N Y Y G S G R Y N W F D P W G Q G T

CGT GAA AAT TAT TAT GGT TCT GGT CGT TAT AAT TGA TTT GAT CCT TGA GGT CAA GGT ACT

CGC GAG AAC TAC TAC GGC TCC GGC CGC TAC AAC TGG TTC GAC CCC TGG GGC CAG GGC ACC

CGA	GGA TCA GGA CGA	CCA	GGA	GGA ACA
CGA	GGA TCA GGA CGA	CCA	GGA	GGA ACA

CGG GGG TCG GGG CGG CCG GGG GGG ACG

AGT

AGC

141

L V T V S S

TTA GTT ACT GTT TCT TCT

TTG GTC ACC GTC TCC TCC

CTT GTA ACA GTA TCA TCA

CTC GTG ACG GTG TCG TCG

CTA AGT AGT

CTG AGC AGC

Oligonucleotides were chemically synthesized based on the base sequence divided for each restriction enzyme-recognition fragment. After sequentially digesting the synthesized oligonucleotide with a corresponding restriction enzyme, a full length of a base sequence encoding the amino acid sequence of the 9F11 antibody variable region by ligation was obtained. cDNA fragments of the 9F11 antibody variable regions of the H-chain and L-chain obtained by the same method with each other (named as rV μ 9F11 and rV κ 9F11, respectively) were integrated into vectors having constant region gene sequences of H-chain and L-chain of the human IgM antibody (C μ and C κ , respectively) by the same method as forming chimera antibodies to obtain recombinant 9F11 μ -chain and κ -chain expression plasmids (rV μ 9F11-C μ and rV κ 9F11-C κ , respectively; Fig. 4).

Example 4. Expression of recombinant antibody

Activities of the antibodies obtained using the plasmids expressing the reconstructed 9F11 antibody genes were investigated with a temporary expression system in COS7 cells (ATCC CRL 1651). Genes were introduced using a mixture of two plasmids (rV μ 9F11-C μ and rV κ 9F11-C κ) and an expression plasmid (Cj) for J-chain of human IgM antibody using a lipofectamine reagent according to the protocol by

GIBCO Co. Cultivation was continued for two days thereafter under a usual culture condition, and the supernatant of the gene-introduced cell culture was retrieved. Recombinant 9F11 antibodies secreted in the culture supernatant were confirmed by subjecting the culture supernatant to an assay system by the sandwich ELISA method using antihuman μ -antibody and antihuman κ -antibody. The antibodies were confirmed to have specificity as described above by FACS analysis of the culture supernatant using cells such as U937/IIIB and MOLT-4/IIIB prepared by infection of U937 cells and MOLT-4 with IIIB strain of HIV-1. Further, the activity of recombinant 9F11 antibody was confirmed by a competitive inhibition test by allowing fluorescence-labeled original 9F11 antigen and the culture supernatant to simultaneously react with U937/IIIB or MOLT-4/IIIB.

Accordingly, it was confirmed that the base sequences of the μ -chain and κ -chain of 9F11 antibody listed in Table 1 are quite important regions responsible for antibody activity against differentiated antigens of the activated lymphocytes expressed in the HIV-infected cells.

It was also confirmed from these results that genes encoding the base sequences of the μ -chain variable region and κ -chain variable region are quite crucial genes for preparing the recombinant anti-HIV antibody and anti-activated lymphocytes.

Industrial Applicability

Since the human IgM monoclonal antibody of the invention against the differentiated antigens expressed in the activated lymphocytes displays a cytolysis function for the activated lymphocytes under mediation by the complement, the antibody can be used as a remedy for controlling abnormally activated lymphocytes in the body. The invention also provides genes encoding the base sequences of the μ -chain variable region and κ -chain variable regions that are quite crucial for preparing the recombinant anti-activated lymphocytes antibody.